AMENDMENTS TO THE CLAIMS

1.-25. (Cancelled)

- 26. (Currently amended) A method for aiding in the determination of whether-a mammal patient is susceptible to or at risk of a disease associated with β-amyloid formation and/or aggregation, said method comprising:
 - (a) determining, in a first sample of brain extract or cerebrospinal fluid obtained from said mammal patient, the amount of a N-terminal truncated and/or post-translationally modified β-amyloid 42 variant selected from the group consisting of Aβ(2-42), Aβ(3-42), Aβ(4-42), Aβ(5-42), Aβ(6-42), Aβ(7-42), Aβ(8-42), and Aβ(9-42);
 - (b) comparing the amount of β-amyloid variant determined in step (a) with the amount of said variant typically present in control samples obtained from one or more patients known to suffer, or known not to suffer[[,]] from said disease associated with β-amyloid formation and/or aggregation;
 - (c) concluding determining, from the comparison in step (b) if the amount of β-amyloid variant determined in step (a) is greater than the amount of said variant typically present in control samples, that whether the mammal patient is susceptible to or at risk of said disease associated with β-amyloid formation and/or aggregation.

27.-34. (Cancelled)

- 35. (Previously presented) The method of claim 26 wherein the post-translationally modified β -amyloid variant is modified by methylation or pyroglutamylation.
- 36. (Previously presented) The method of claim 35 wherein the methylation is present at position 1, 2, 4, or 6 of an N-terminal truncated β-amyloid variant.
- 37. (Withdrawn) The method according to claim 35 further characterized in that the pyroglutamylation is present at position 3 of an N-terminal truncated β -amyloid variant starting at position 3 of β -amyloid.
- 38. (Cancelled)
- 39. (Cancelled)

- 40. (Currently amended) The method of claim 26 wherein at least one of the first and second samples the sample is a brain extract sample or a body fluid sample.
- 41. (Currently amended) The method of claim <u>26</u> [[40]] wherein the body fluid sample is a blood sample or a cerebrospinal fluid (CSF) sample.
- 42. (Previously presented) The method of claim 26 wherein the disease associated with β -amyloid formation and/or aggregation is Alzheimer's disease (AD).
- 43. (Currently amended) The method of claim 42 [[26]] wherein the susceptibility to Alzheimer's disease (AD) or the risk of developing AD is determined by detecting A β (5-42) or A β (8-42) in a body fluid sample obtained from the mammal.
- 44.-56. (Cancelled)
- 57. (Currently amended) The method of claim [[56]] $\underline{26}$ wherein said β -amyloid variant is A β (4-42).
- 58. (Previously presented) The method of claim 26 wherein the post-translationally modified β-amyloid variant is modified by methylation.
- 59. (Previously presented) The method of claim 58 wherein the methylation is present at position 4 of an N-terminal truncated β-amyloid variant.
- 60. (Currently amended) The method of claim $\underline{42}$ [[26]] wherein the susceptibility to Alzheimer's disease (AD) or the risk of developing AD is determined by detecting A β (5-42) in a body fluid sample obtained from the mammal.
- 61. (Previously presented) The method of claim 26 wherein the amount of N-terminal truncated and/or post-translationally modified β-amyloid variant is determined by 2-D electrophoresis or mass spectrometry or both.
- 62. (Cancelled)

- 63. (Previously presented) The method of claim 26 wherein the amount of the N-terminal truncated and/or post-translationally modified β -amyloid 42 ($A\beta_{42}$) variant is detected using an antibody that binds an epitope at the N-terminus of said variant.
- 64. (New) The method of claim 26 wherein the disease associated with β-amyloid formation and/or aggregation is mild cognitive impairment (MCI) progressing to Alzheimer's disease.